Roberto Malinow *et al.* Serial No. 09/353,126

## **Amendments** to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application. No amendments to the claims have been made at this time.

## **Listing of Claims:**

- 1.-12. (cancelled)
- 13. (previously presented) An in vitro method for screening for candidate drugs for the treatment of Alzheimer's disease, said method comprising:

contacting slices of mouse hippocampal tissue containing cells, having a PS-1  $\Delta$ 9 mutation and having enhanced synaptic potentiation upon stimulation as compared to wild-type hippocampal cells with a candidate drug;

subjecting said mutant hippocampal cells to tetanic stimulation; and determining the effect of said candidate drug on the synaptic potentiation of said mutant hippocampal cells;

wherein a reduction in the enhanced synaptic potentiation of the mutant hippocampal cells is indicative of activity of a candidate drug for the treatment of Alzheimer's disease.

14. (previously presented) An in vitro method for screening for candidate drugs for the treatment of Alzheimer's disease, said method comprising:

contacting mammalian hippocampal cells comprising a PS-1 Δ9 presentilin gene mutation wherein said hippocampal cells have enhanced synaptic potentiation upon stimulation as compared to wild-type hippocampal cells with a candidate drug;

subjecting said mutant hippocampal cells to tetanic stimulation; and determining the effect of said candidate drug on the synaptic potentiation of said mutant hippocampal cells;

wherein a reduction in the enhanced synaptic potentiation of the mutant hippocampal cells is indicative of activity of a candidate drug for the treatment of

Roberto Malinow *et al.* Serial No. 09/353,126

Alzheimer's disease.

- 15. (previously presented) The method according to Claim 14, wherein mouse hippocampal tissue slices comprise said mutant hippocampal cells.
- 16. (previously presented) The method according to Claim 14, wherein said enhanced synaptic potentiation is a result of a change in the GABA<sub>A</sub> receptor pathway.
- 17. (previously presented) An in vitro method for screening for candidate drugs for the treatment of Alzheimer's disease, said method comprising:

contacting mammalian hippocampal cells comprising a PS-1  $\Delta$ 9 presenting gene mutation and having enhanced synaptic potentiation upon stimulation as compared to wild-type hippocampal cells with a candidate drug;

subjecting said mutant hippocampal cells and said wild-type hippocampal cells to a tetanic stimulus;

measuring changes in potentiation with time of the mutant hippocampal cells and wild-type hippocampal cells and comparing the effect of said candidate drug on the synaptic potentiation of said mutant hippocampal cells as compared to the observed synaptic potentiation of said wild-type hippocampal cells;

wherein a reduction in the enhanced synaptic potentiation of the mutant hippocampal cells as compared to the synaptic potentiation of the wild-type cells is indicative of activity of a candidate drug for the treatment of Alzheimer's disease.

18.-19. (cancelled)

20. (previously presented) An in vitro method for screening for candidate drugs for the treatment of Alzheimer's disease, said method comprising:

contacting mammalian hippocampal cells comprising a PS-1  $\Delta 9$  presentlin gene

Roberto Malinow *et al.* Serial No. 09/353,126

mutation and having enhanced synaptic potentiation upon stimulation as compared to wild-type hippocampal cells with a candidate drug;

subjecting said mutant hippocampal cells and said wild-type hippocampal cells to a tetanic stimulus at a first potential of glutamate currents and a second potential of GABA<sub>A</sub> currents;

measuring the synaptic response at each of the first and second potentials for said mutant hippocampal cells and said wild-type hippocampal cells and comparing the effect of said candidate drug on said mutant hippocampal cells and said wild-type hippocampal cells;

wherein a reduction in the enhanced synaptic response of the mutant hippocampal cells without a significant change in the synaptic response of the wild-type cells is indicative of activity of a candidate drug for the treatment of Alzheimer's disease.

21. (previously presented) An in vitro method for screening for candidate drugs for the treatment of Alzheimer's disease, said method comprising:

contacting mouse hippocampal cells comprising a PS-1  $\Delta$ 9 presentiin-1 gene mutation and having enhanced synaptic potentiation upon tetanic stimulation as compared to wild-type hippocampal cells, with a candidate drug;

subjecting said mutant hippocampal cells and said wild-type hippocampal cells to tetanic stimulation; and

comparing the effect of said candidate drug on said mutant hippocampal cells and said wild-type hippocampal cells upon tetanic stimulation;

wherein a reduction in the enhanced synaptic potentiation of the mutant hippocampal cells without a significant change in the synaptic potentiation of the wild-type cells is indicative of activity of a candidate drug for the treatment of Alzheimer's disease.

## 22.-25. (cancelled)

Roberto Malinow et al. Serial No. 09/353,126

26. (previously presented) A method for screening for a candidate drug that suppresses intracellular calcium rise in slices of mouse hippocampal tissue containing cells having a PS-1  $\Delta$ 9 mutation in a presentilin gene combined with a candidate drug for the treatment of Alzheimer's disease, said method comprising:

contacting hippocampal cells comprising a presentilin gene mutation and having enhanced synaptic potentiation upon stimulation as compared to wild-type hippocampal cells with a candidate drug that suppresses intracellular calcium rise in said cells; subjecting said mutant hippocampal cells to tetanic stimulation; and

determining the effect of said candidate drug on the ratio of peak inhibitory to excitory responses;

wherein an enhanced said ratio of peak inhibitory to excitory responses in said mutant hippocampal cells as compared to wild-type hippocampal cells is indicative of activity of a candidate drug for the treatment of Alzheimer's disease.